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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/903,376	07/10/2001	Thomas J. Brennan	R-599	8327
7590 11/22/2002 DELTAGEN, INC.			EXAMINER	
1003 Hamilton Avenue Menlo Park, CA 94025			PARAS JR, PETER	
14104110 2 41-11, 1-1			ART UNIT	PAPER NUMBER
			1632 DATE MAILED: 11/22/2002	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)
	09/903,376	BRENNAN, THOMAS J.
Office Action Summary	Examiner	Art Unit
	Peter Paras, Jr.	1632
The MAILING DATE of this communication	n appears on the cover sheet w	vith the correspondence address
Period for Reply		
A SHORTENED STATUTORY PERIOD FOR R THE MAILING DATE OF THIS COMMUNICATI - Extensions of time may be available under the provisions of 37 C after SIX (6) MONTHS from the mailing date of this communication - If the period for reply specified above is less than thirty (30) days - If NO period for reply is specified above, the maximum statutory properties to reply within the set or extended period for reply will, by - Any reply received by the Office later than three months after the earned patent term adjustment. See 37 CFR 1.704(b). Status	ION. CFR 1.136(a). In no event, however, may a con. s, a reply within the statutory minimum of the period will apply and will expire SIX (6) MC a statute, cause the application to become A	a reply be timely filed nirty (30) days will be considered timely. DNTHS from the mailing date of this communication. ABANDONED (35 U.S.C. § 133).
1) Responsive to communication(s) filed or	ı	
2a) This action is FINAL . 2b) ⊠	This action is non-final.	
3) Since this application is in condition for a	allowance except for formal m	atters, prosecution as to the merits is
closed in accordance with the practice u Disposition of Claims	nder Ex parte Quayle, 1935 C	;.D. 11, 453 O.G. 213.
4)⊠ Claim(s) <u>1-27</u> is/are pending in the applic	cation.	
4a) Of the above claim(s) is/are wit	hdrawn from consideration.	
5) Claim(s) is/are allowed.		
6) Claim(s) is/are rejected.		
7) Claim(s) is/are objected to.	•	
8) Claim(s) <u>1-27</u> are subject to restriction an	d/or election requirement.	
Application Papers		
9) The specification is objected to by the Exa	<u></u>	. Also Espaining
10) The drawing(s) filed on is/are: a)		
Applicant may not request that any objection		
11) The proposed drawing correction filed on _ If approved, corrected drawings are required		disapproved by the Examiner.
12) The oath or declaration is objected to by the		
Priority under 35 U.S.C. §§ 119 and 120	TO Examinor.	
13) Acknowledgment is made of a claim for for	oreian priority under 35 H S C	& 119(a)-(d) or (f)
a) ☐ All b) ☐ Some * c) ☐ None of:	oreign priority under 55 0.6.0	. 3 113(a)-(a) or (i).
1. Certified copies of the priority docu	ments have been received	
Certified copies of the priority docu		Application No.
Copies of the certified copies of the		
application from the Internation * See the attached detailed Office action for	al Bureau (PCT Rule 17.2(a))).
14) Acknowledgment is made of a claim for do	mestic priority under 35 U.S.C	. § 119(e) (to a provisional application).
a) ☐ The translation of the foreign languages 15)☐ Acknowledgment is made of a claim for do	•	
Attachment(s)	•	* *
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-94 3) Information Disclosure Statement(s) (PTO-1449) Paper N	18) 5) Notice o	w Summary (PTO-413) Paper No(s) of Informal Patent Application (PTO-152)

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DETAILED ACTION

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claims 1-4, drawn to a targeting construct comprising nucleotide sequences homologous to a 5-HT-2B gene and a method of producing a targeting construct, classified in class 435, subclass 320.1.
- II. Claims 5-7, 9, and 24 drawn to cells comprising a disruption in a 5-HT-2B gene, classified in class 435, subclass 325.
- III. Claims 8, 10, and 17-23, drawn to a transgenic non-human animal, particularly a mouse, comprising a disruption in a 5-HT-2B gene, and a method of making the same, classified in classes 800, 800, and 800 subclass 13, 18, and 25.
- IV. Claims 11-12, drawn to methods of identifying agents that modulate the expression of a 5-HT-2B gene or modulate the function of a 5-HT-2B comprising screening said agents in a transgenic non-human animal, classified in class 800, subclass 3.
- V. Claims 13-15, drawn to methods of identifying agents that modulate expression of a 5-HT-2B gene or function of a 5-HT-2B in a cell *in vitro*, classified in class 435, subclass 7.2.
- VI. Claim 16, drawn to an unknown agent is unclassifiable.

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VII. Claim 27, drawn to phenotypic data, in an electronic database, associated with a transgenic mouse, classified in class 702, subclass 19.

- VIII. Claim 25, drawn to methods of identifying an agent that modulates a phenotype associated with or behavior associated with a disruption in a 5-HT-2B gene, comprising screening agents in a transgenic mouse, classified in class 800, subclass 3.
- IX. Claim 26, drawn to an agonist or antagonist of a 5-HT-2B receptor is unclassifiable as the agonist or antagonist is unknown.

The products of Inventions I, II, III, VI, VII, and IX each from the other are distinct each from the other. Inventions are distinct if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions have different modes of operation, different function, and different effects. The products of Groups I, II, III, VI, VII, and IX have different chemical structures, are made by different methods, and can be used in different methods which require different technical considerations and materially different reagents. For example, the transgenic animal non-human animal of Group III can be used as a model of disease while the targeting construct of Group I may be used to disrupt a gene in a somatic cell *in vitro*, the cells of Group II may be used to isolate a protein, and the data of Group VII may be used for statistical analysis in a database. Also, the agent of group VI has a different chemical structure from the targeting construct, cells, and transgenic

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non-human animals of Groups I, II, and III respectively, and may be used in different methods, which require different technical considerations with respect to modulation of a 5-HT-2B. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, different classifications, and separate search requirement, restriction for examination purposes as indicated is proper.

Although there are no provisions under the section for "Relationship of Inventions" in MPEP 806.05 for inventive groups that are directed to different methods, restriction is deemed to be proper between groups IV, V, and VIII because their methods appear to constitute patentably distinct inventions, each with a distinct purpose and further comprising distinct methodologies and using different products. For example, the method of Group IV requires the use of a transgenic non-human animal while the method of Group V requires the use of a cell *in vitro*. Because these inventions are distinct for the reasons given above and a separate search is required for each of Groups IV, V and VIII, restriction for examination purposes as indicated is proper.

The products of Inventions I, II, III, VI, VII, and IX and the methods of Invention IV, V, VIII are distinct. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions have different modes of operation, different function, and different effects each from the other. The products of Groups I, II, III, VI, VII, and IX can

be used in methods that require different technical considerations and materially different reagents from the methods of Groups IV, V, VIII. The method of Group IV can be practiced with products that have different chemical structures than the products of Groups I, II, III, VI, VII, and IX. For example, the transgenic animals of Group III may be used to produce antibodies while the method of Group IV may be used to identify agents that modulate the expression of a 5-HT-2B. Further, the method of Group IV may be practiced with agents that have different chemical structures from the agent of Group VI. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, different classifications, and separate search requirement, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Any inquiry concerning this communication or earlier communications from the examiner(s) should be directed to Peter Paras, Jr., whose telephone number is 703-308-8340. The examiner can normally be reached Monday-Friday from 8:30 to 4:30 (Eastern time).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Deborah Reynolds, can be reached at 703-305-4051. Papers related to this application may be submitted by facsimile transmission. Papers should be faxed via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The CM1 Fax Center numbers are (703) 308-4242 and (703) 305-3014.

Inquiries of a general nature or relating to the status of the application should be directed to Dianiece Jacobs whose telephone number is (703) 305-3388.

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Peter Paras, Jr.

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